aminopropazine fumarate equivalent to 25 milligrams of aminopropazine base.

- (b) *Sponsor*. See No. 000061 in §510.600(c) of this chapter.
- (c) *Conditions of use.* (1) The drug is used for reducing excessive smooth muscle contractions, such as occur in urethral spasms associated with urolithiasis in cats and dogs and in colic spasms in horses.¹
- (2) It is administered intramuscularly or intravenously to dogs and cats at a level of 1 to 2 milligrams per pound of body weight. It is administered intramuscularly or intravenously to horses at a level of 0.25 milligrams per pound of body weight. Dosage can be repeated every 12 hours, as indicated.1
- (3) Not for use in animals intended for food purposes.¹
- (4) For use only by or on the order of a licensed veterinarian.¹

[40 FR 13858, Mar. 27, 1975, as amended at 46 FR 48642, Oct. 2, 1981; 61 FR 8873, Mar. 6, 1996; 62 FR 61625, Nov. 19, 1997]

§ 522.84 Beta-aminopropionitrile fumarate.

- (a) Specifications. Each vial contains 7.0 milligrams of beta-aminopropionitrile fumarate sterile lyophilized powder which is reconstituted for injection with 10 milliliters of sterile physiologic saline, USP.
- (b) Sponsor. See No. 064146 in \$510.600(c) of this chapter.
 - (c) [Reserved]
- (d) Conditions of use—(1) Horses—(i) Amount. 7 milligrams (10 milliliters) intralesionally every other day for 5 treatments beginning about 30 days after initial injury.
- (ii) *Indications for use.* For treatment of tendinitis of the superficial digital flexor tendon (SDFT) in the adult horse where there is sonographic evidence of fiber tearing.
- (iii) *Limitations.* Single dose container for intralesional injection. Do not use in horses with dermal irritation or open skin lesions in the injection area. Do not administer

intraarticularly, into the tendon sheath, or in the presence of concurrent limb fractures. Do not use in breeding animals since the effects on fertility, pregnancy, or fetal health have not been determined. Not for use in horses intended for food. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

(2) [Reserved]

[63 FR 44382, Aug. 19, 1998]

§522.88 Sterile amoxicillin trihydrate for suspension.

- (a)(1) Specifications. Each vial contains 3 grams of amoxicillin as the trihydrate. The powder is reconstituted with sterile water for injection USP to a concentration of 100 or 250 milligrams per milliliter for use as in paragraph (d) of this section.
- (2) Each vial contains 25 grams of amoxicillin as the trihydrate. The powder is reconstituted with sterile water for injection USP to a concentration of 250 milligrams per milliliter for use as in paragraph (e).
- (b) Sponsor. See 000069 in \$510.600(c) of this chapter.
- (c) Related tolerance. See §556.38 of this chapter.
- (d) *Conditions of use in dogs and cats—*(1) *Amount.* 5 milligrams per pound of body weight daily.
- (2) Indications for use—(i) Dogs. Treatment of infections caused by susceptible strains of organisms as follows: Respiratory infections (tonsillitis, tracheobronchitis) due to Staphylococcus aureus, Streptococcus spp., Escherichia coli, and Proteus mirabilis; genitourinary infections (cystitis) due to S. aureus, Streptococcus spp., E. coli, and P. $\it mirabilis$; gastrointestinal infections (bacterial gastroenteritis) due to $\it S$. aureus, Streptococcus spp., E. coli, and P. mirabilis; bacterial dermatitis due to S. scesses, lacerations, and wounds), due to S. aureus, Streptococcus spp., E. coli, and P. mirabilis.
- (ii) Cats. Treatment of infections caused by susceptible strains of organisms as follows: Upper respiratory infections due to S. aureus, Staphylococcus spp., Streptococcus spp., Hemophilus spp., E. coli, Pasteurella spp.,

¹These conditions are NAS/NRC reviewed and deemed effective. Applications for these uses need not include effectiveness data as specified by §514.111 of this chapter, but may require bioequivalency and safety information.

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and *P. mirabilis*; genitourinary infections (cystitis) due to *S. aureus, Streptococcus* spp., *E. coli, P. mirabilis*, and *Corynebacterium* spp.; gastrointestinal infections due to *E. coli, Proteus* spp., *Staphylococcus* spp., and *Streptococcus* spp.; skin and soft tissue infections (abscesses, lacerations, and wounds) due to *S. aureus, Staphylococcus* spp., *Streptococcus* spp., *E. coli*, and *Pasteurella multocida*.

- (3) Limitations. For use in dogs and cats only. Administer once daily for up to 5 days by intramuscular or subcutaneous injection. Continue treatment for 48 hours after the animal has become afebrile or asymptomatic. If no improvement is seen within 5 days, review the diagnosis and change therapy. As with all antibiotics, appropriate in vitro culturing susceptibility testing of samples taken before treatment should be conducted. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
- (e) *Condition of use. Cattle*—(1) *Amount.* 3 to 5 milligrams per pound of body weight once a day according to the animal being treated, the severity of infection, and the animal's response.
- (2) Indications for use.— Treatment of diseases due to amoxicillin-susceptible organisms as follows: Respiratory tract infections (shipping fever, pneumonia) due to *P. multocida, P. hemolytica, Hemophilus* spp., Staphylococcus spp., and Streptococcus spp. and acute necrotic pododermatitis (foot rot) due to Fusobacterium necrophorum.
- (3) Limitations. Administer once daily for up to 5 days by intramuscular or subcutaneous injection. Continue treatment for 48 to 72 hours after the animal has become afebrile or asymptomatic. Do not continue treatment beyond 5 days. Treated animals must not be slaughtered for food during treatment and for 25 days after the last treatment. As with all antibiotics, appropriate in vitro culturing and susceptibility testing of samples taken before treatment should be conducted. Milk from treated cows must not be used for human consumption during treatment or for 96 hours (8 milkings) after last treatment. Maximum volume per injection should not exceed 30 milliliters. Federal law restricts this drug to use

by or on the order of a licensed veterinarian.

[57 FR 37330, Aug. 18, 1992; 60 FR 55659, Nov. 2, 1995]

§ 522.90 Ampicillin implantation and injectible dosage forms.

§ 522.90a Ampicillin trihydrate sterile suspension.

- (a) *Specifications*. Each milliliter contains ampicillin trihydrate equivalent to 200 milligrams of ampicillin.
- (1) *Sponsor.* See No. 053501 in §510.600(c) of this chapter.
- (2) Related tolerances. See §556.40 of this chapter.
 - (3) Conditions of use—(i) Calves.
- (A) Amount. For enteritis: 3 milligrams per pound of body weight, intramuscularly, once or twice daily, for up to 3 days. For pneumonia: 3 milligrams per pound of body weight, intramuscularly, twice daily, for up to 3 days.
- (B) *Indications for use.* Treatment of bacterial enteritis caused by *Escherichia coli* and bacterial pneumonia caused by *Pasteurella* spp. susceptible to ampicillin.
- (C) Limitations. Not for use in other animals raised for food production. Treated animals must not be slaughtered for food use during treatment or for 9 days after the last treatment. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
- (ii) *Dogs.* (A) *Amount.* 3 to 6 milligrams per pound of body weight intramuscularly, once or twice daily.
- (B) Indications for use. Treatment of respiratory tract infections due to E. coli, Pseudomonas spp., Proteus spp., Staphylococcus spp., and Streptococcus spp.; tonsillitis due to E. coli, Pseudomonas spp., Streptococcus spp., and Staphylococcus spp.; generalized infections (septicemia) associated with abscesses, lacerations, and wounds due to Staphylococcus spp. and Streptococcus spp.
- (C) Limitations. Continue treatment at least 48 hours after the animal's temperature has returned to normal and other signs of infection have subsided. Usual treatment is 3 to 5 days. Federal law restricts this drug to use